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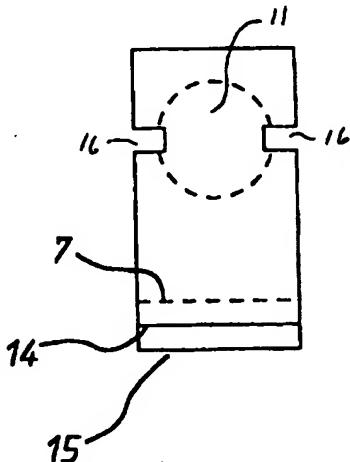
INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6 : G01N 27/403		A1	(11) International Publication Number: WO 97/18464 (43) International Publication Date: 22 May 1997 (22.05.97)
(21) International Application Number: PCT/AU96/00724		(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN. ARIPO patent (KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).	
(22) International Filing Date: 15 November 1996 (15.11.96)			
(30) Priority Data: PN 6619 16 November 1995 (16.11.95) AU			
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(54) Title: ELECTROCHEMICAL CELL

(57) Abstract

A method of manufacture of a thin layer electrochemical cell (Figs 12, 14) comprising the steps of: forming an aperture (11) extending through a sheet (1) of electrically resistive material, said aperture defining a side wall of the cell; mounting a first thin electrode layer (13) to one side of the sheet and extending over aperture (11) whereby to define a cell first end wall; mounting a second thin electrode layer (13) to the other side of the sheet and extending over aperture (11) whereby to define a second cell end wall in substantial overlying registration with the first electrode; and providing means (16) for admission of a liquid into the cell.



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TITLE: ELECTROCHEMICAL CELL**FIELD OF THE INVENTION**

This invention relates to an electrochemical cell for determining the concentration of an analyte in a carrier.

5 BACKGROUND ART

The invention herein described is an improvement in or modification of the invention described in our co-pending application PCT/AU96/00365, the contents of which are incorporated herein by reference.

The invention will herein be described with particular reference to a biosensor 10 adapted to measure the concentration of glucose in blood, but it will be understood not to be limited to that particular use and is applicable to other analytic determinations.

It is known to measure the concentration of a component to be analysed in an aqueous liquid sample by placing the sample into a reaction zone in an electrochemical cell comprising two electrodes having an impedance which renders them suitable for 15 amperometric measurement. The component to be analysed is allowed to react directly or indirectly with a redox reagent whereby to form an oxidisable (or reducible) substance

- 2 -

in an amount corresponding to the concentration of the component to be analysed. The quantity of the oxidisable (or reducible) substance present is then estimated electrochemically. Generally this method requires sufficient separation of the electrodes so that electrolysis products at one electrode cannot reach the other electrode and

5 interfere with the processes at the other electrode during the period of measurement.

In our co-pending application we described a novel method for determining the concentration of the reduced (or oxidised) form of a redox species in an electrochemical cell of the kind comprising a working electrode and a counter (or counter/reference) electrode spaced from the working electrode by a predetermined distance. The method

10 involves applying an electric potential difference between the electrodes and selecting the potential of the working electrode such that the rate of electro-oxidation of the reduced form of the species (or of electro-reduction of the oxidised form) is diffusion controlled. The spacing between the working electrode and the counter electrode is selected so that reaction products from the counter electrode arrive at the working

15 electrode. By determining the current as a function of time after application of the potential and prior to achievement of a steady state current and then estimating the magnitude of the steady state current, the method previously described allows the diffusion coefficient and/or the concentration of the reduced (or oxidised) form of the species to be estimated.

20 Our co-pending application exemplifies this method with reference to use of a "thin layer electrochemical cell" employing a GOD/Ferrocyanide system. As herein used, the term "thin layer electrochemical cell" refers to a cell having closely spaced electrodes such that reaction product from the counter electrode arrives at the working

- 3 -

electrode. In practice, the separation of electrodes in such a cell for measuring glucose in blood will be less than 500 microns, and preferably less than 200 microns.

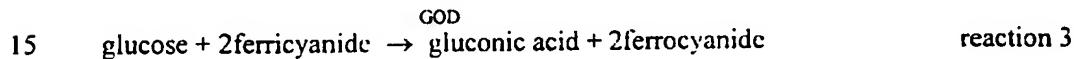
The chemistry used in the exemplified electrochemical cell is as follows:



where GOD is the enzyme glucose oxidase, and GOD* is the 'activated' enzyme.

Ferricyanide ($[\text{Fe}(\text{CN})_6]^{3-}$) is the 'mediator' which returns the GOD* to its catalytic state. GOD, an enzyme catalyst, is not consumed during the reaction so long as excess mediator is present. Ferrocyanide ($[\text{Fe}(\text{CN})_6]^{4-}$) is the product of the total reaction.

10 Ideally there is initially no ferrocyanide, although in practice there is often a small quantity. After reaction is complete the concentration of ferrocyanide (measured electrochemically) indicates the initial concentration of glucose. The total reaction is the sum of reactions 1 and 2:



"Glucose" refers specifically to β -D-glucose.

The prior art suffers from a number of disadvantages. Firstly, sample size required is greater than desirable. It would be generally preferable to be able to make measurements on samples of reduced volume since this in turn enables use of less invasive methods to obtain samples.

Secondly, it would be generally desirable to improve the accuracy of measurement and to eliminate or reduce variations due, for example, to cell asymmetry or other factors introduced during mass production of microcells. Likewise, it would be desirable to reduce electrode "edge" effects.

Thirdly, since the cells are disposable after use, it is desirable that they be capable of mass production at relatively low cost.

DISCLOSURE OF THE INVENTION

According to one aspect the invention consists in a method of manufacture of

5 an electrochemical cell comprising the steps of:

forming an aperture extending through a sheet of electrically resistive material,

said aperture defining a side wall of the cell;

mounting a first thin electrode layer to one side of the sheet and extending over the aperture whereby to define a cell first end wall;

10 mounting a second thin electrode layer to the other side of the sheet and

extending over the aperture whereby to define a cell second end wall in substantial

overlying registration with the first electrode; and

providing means for admission of a liquid into the cell defined between the side wall and said end walls.

15 The first and second electrode layers may be conductors or semi-conductors and may be the same or different. Noble metal electrode layers are preferred.

In preferred embodiments of the invention the aperture is of circular cross-section whereby the side wall is cylindrical and the first and second electrodes cover the aperture.

20 DESCRIPTION OF PREFERRED EMBODIMENTS

The invention will now be particularly described by way of example only with reference to the accompanying schematic drawings wherein:

Figure 1 shows the product of manufacturing step 2 in plan.

Figure 2 shows the product of Figure 1 in side elevation.

- 5 -

Figure 3 shows the product of Figure 1 in end elevation.

Figure 4 shows the product of manufacturing step 3 in plan.

Figure 5 shows the product of Figure 4 in cross-section on line 5-5 of Figure 4.

Figure 6 shows the product of manufacturing step 5 in plan.

5 Figure 7 shows the product of Figure 6 in side elevation.

Figure 8 shows the product of Figure 6 in end elevation.

Figure 9 shows the product of manufacturing step 7 in plan.

Figure 10 is a cross-section of Figure 9 on line 10-10.

Figure 11 shows the product of Figure 9 in end elevation.

10 Figure 12 shows a cell according to the invention in plan.

Figure 13 shows the cell of Figure 12 in side elevation.

Figure 14 shows the cell of Figure 12 in end elevation.

Figure 15 shows a scrap portion of a second embodiment of the invention in
enlarged section.

15 The construction of a thin layer electrochemical cell will now be described by
way of example of the improved method of manufacture.

Step 1: A sheet 1 of Melinex® (a chemically inert, and electrically resistive
Polyethylene Terephthalate ["PET"]) approximately 13cm x 30cm and 100 micron thick
was laid flat on a sheet of release paper 2 and coated using a Number 2 MYAR bar to a
20 thickness of 12 microns wet (approximately 2-5 microns dry) with a water-based heat
activated adhesive 3 (ICI Novacoat system using catalyst:adhesive). The water was then
evaporated by means of a hot air dryer leaving a contact adhesive surface. The sheet was
then turned over on a release paper and the reverse side was similarly coated with the
same adhesive 4, dried, and a protective release paper 5 applied to the exposed adhesive

surface. The edges were trimmed to obtain a sheet uniformly coated on both sides with tacky contact adhesive protected by release paper.

Step 2: The sheet with protective release papers was cut into strips 7, each about 18mm x 210mm (Figures 1-3).

5 Step 3: A strip 7 of adhesive-coated PET from step 2 with release paper 2, 5 on respective sides, was placed in a die assembly (not shown) and clamped. The die assembly was adapted to punch the strip with a locating hole 10 at each end and with for example 37 circular holes 11 each of 3.4mm diameter at 5mm centres equi-spaced along a line between locating holes 10. The area of each hole 11 is approximately 9 square
10 mm.

Step 4: A sheet 12 of Mylar[®] PET approximately 21cm square and 135 microns thick was placed in a sputter coating chamber for palladium coating 13. The sputter coating took place under a vacuum of between 4 and 6 millibars and in an atmosphere of argon gas. Palladium was coated on the PET to a thickness of 100-1000
15 angstroms. There is thus formed a sheet 14 having a palladium sputter coating 13.

Step 5: The palladium coated PET sheet 14 from Step 4 was then cut into strips 14 and 15 and a die was used to punch two location holes 16 in each strip, at one end (Figures 6, 7 and 8). Strips 14 and 15 differ only in dimension strips 14 being 25mm x 210mm and strips 15 being 23mm x 210mm.

20 Step 6: A spacer strip 7 prepared as in step 3 was then placed in a jig (not shown) having two locating pins (one corresponding to each locating hole 10 of strip 7) and the upper release paper 2 was removed. A strip 14 of palladium coated PET prepared as in step 5 was then laid over the adhesive layer, palladium surface downwards, using the jig pins to align the locating holes 16 with the underlying PET

strip 7. This combination was then passed through a laminator comprising a set of pinch rollers, one of which was adapted to heat the side bearing a palladium coated PET strip 14. The roller on the opposite side of the strip 7 was cooled. By this means, only the adhesive between the palladium of strip 14 and PET strip 7 was activated.

5 Step 7: PET strip 7 was then turned over and located in the jig with the release coating uppermost. The release coating was peeled off and second palladium coated strip 15 was placed palladium side down on the exposed adhesive surface using the locating pins to align the strips. this assembly was now passed again through the laminator of step 6, this time with the hot roll adjacent the palladium coated Mylar[®] added in step 7 so as to activate the intervening adhesive (Figures 9, 10 and 11).

10 Step 8: The assembly from step 7 was returned to the die assembly and notches 17 punched in locations so as to extend between the circular holes 11 previously punched in the Melinex[®] PET and the strip edge 17. Notches 16 extend so as to intercept the circumference of each circular cell. The strip was then guillotined to give 15 37 individual "sensor strips", each strip being about 5mm wide and each having one thin layer cavity cell (Figures 12, 13 and 14).

17 There is thus produced a cell as shown in Figures 12, 13 or 14. The cell comprises a first electrode consisting of PET layer 12, a palladium layer 13, an adhesive layer 3, a PET sheet 1, a second adhesive layer 4, a second electrode comprising 20 palladium layer 13, and a PET layer 12. Sheet 1 defines a cylindrical cell 11 having a thickness in the cell axial direction corresponding to the thickness of the Melinex[®] PET sheet layer 1 together with the thickness of adhesive layers 3 and 4. The cell has circular palladium end walls. Access to the cell is provided at the side edge of the cell where notches 16 intersect cell 11.

In preferred embodiments of the invention, a sample to be analysed is introduced to the cell by capillary action. The sample is placed on contact with notch 16 and is spontaneously drawn by capillary action into the cell, displaced air from the cell venting from the opposite notch 16. A surfactant may be included in the capillary space

5 to assist in drawing in the sample.

The sensors are provided with connection means for example edge connectors whereby the sensors may be placed into a measuring circuit. In a preferred embodiment this is achieved by making spacer 1 shorter than palladium supporting sheets 14, 15 and by making one sheet 15 of shorter length than the other 14. This forms a socket region

10 20 having contact areas 21, 22 electrically connected with the working and counter electrodes respectively. A simple tongue plug having corresponding engaging conduct surfaces can then be used for electrical connection. Connectors of other form may be devised.

Chemicals for use in the cell may be supported on the cell electrodes or walls,

15 may be supported on an independent support contained within the cell or may be self-supporting.

In one embodiment, chemicals for use in the cell are printed onto the palladium surface of the electrode immediately after step 1 at which stage the freshly-deposited palladium is more hydrophilic. For example, a solution containing 0.2 molar potassium ferricyanide and 1% by weight of glucose oxidase dehydrogenase may be printed on to

20 the palladium surface. Desirably, the chemicals are printed only in the areas which will form a wall of the cell and for preference the chemicals are printed on the surface by means of an ink jet printer. In this manner, the deposition of chemicals may be precisely controlled. If desired, chemicals which are desirably separated until required for use

may be printed respectively on the first and second electrodes. For example, a GOD/ferrocyanide composition can be printed on one electrode and a buffer on the other. Although it is highly preferred to apply the chemicals to the electrodes prior to assembly into a cell, chemicals may also be introduced into the cell as a solution after 5 step 6 or step 8 by pipette in the traditional manner and the solvent subsequently is removed by evaporation or drying. Chemicals need not be printed on the cell wall or the electrodes and may instead be impregnated into a gauze, membrane, non-woven fabric or the like contained within, or filling, the cavity (eg inserted in cell 11 prior to steps 6 or 7). In another embodiment the chemicals are formed into a porous mass which may be 10 introduced into the cell as a pellet or granules. Alternatively, the chemicals may be introduced as a gel.

In a second embodiment of the invention a laminate 21 is first made from a strip 14 as obtained in step 5 adhesively sandwiched between two strips 7 as obtained from step 3. Laminate 20 is substituted for sheet 1 in step 5 and assembled with electrodes as 15 in steps 6 and 7.

There is thus obtained a cell as shown in Figure 15 which differs from that of Figures 9 to 11 in that the cell has an annular electrode disposed between the first and second electrode. This electrode can for example be used as a reference electrode.

It will be understood that in mass production of the cell, the parts may be 20 assembled as a laminate on a continuous line. For example, a continuous sheet 1 of PET could be first punched and then adhesive could be applied continuously by printing on the remaining sheet. Electrodes (pre-printed with chemical solution and dried) could be fed directly as a laminate onto the adhesive coated side. Adhesive could then be applied

to the other side of the punched core sheet and then the electrode could be fed as a laminate onto the second side.

The adhesive could be applied as a hot melt interleaving film. Alternatively, the core sheet could first be adhesive coated and then punched.

5 By drying chemicals on each electrode prior to the gluing step the electrode surface is protected from contamination.

Although the cell has been described with reference to Mylar® and Melinex® PET, other chemically inert and electrically resistive materials may be utilised and other dimensions chosen. The materials used for spacer sheet 1 and for supporting the
10 reference and counter electrodes may be the same or may differ one from the other.

Although the invention has been described with reference to palladium electrodes, other metals such as platinum, silver, gold, copper or the like may be used and silver may be reacted with a chloride to form a silver/silver chloride electrode or with other halides.

The electrodes need not be of the same metal.

15 Although the use of heat activated adhesives has been described, the parts may be assembled by use of hot melt adhesives, fusible laminates and other methods.

The dimensions of the sensor may readily be varied according to requirements.

While it is greatly preferred that the electrodes cover the cell end openings, in other embodiments (not illustrated) the electrodes do not entirely cover the cell end
20 openings. In that case it is desirable that the electrodes be in substantial overlying registration.

Preferred forms of the invention in which the electrodes cover the apertures of cell 11 have the advantages that the electrode area is precisely defined simply by

punching hole 11. Furthermore the electrodes so provided are parallel, overlying, of substantially the same area, and are substantially or entirely devoid of "edge" effects.

Although in the embodiments described each sensor has one cell cavity, sensors may be provided with two or more cavities. For example, a second cavity may be 5 provided with a predetermined quantity of the analyte and may function as a reference cell.

As will be apparent to those skilled in the art from the teaching herein contained, a feature of one embodiment herein described may be combined with features of other embodiments herein described or with other embodiments described in our co-pending application. Although the sensor has been described with reference to 10 palladium electrodes and a GOD/ferrocyanide chemistry, it will be apparent to those skilled in the art that other chemistries, and other materials of construction may be employed without departing from the principles herein taught.

CLAIMS:-

1. A method of manufacture of a thin layer electrochemical cell (as herein defined) comprising the steps of:

forming an aperture extending through a sheet of electrically resistive material,

5 said aperture defining a side wall of the cell,

mounting a first thin electrode layer to one side of the sheet and extending over the aperture whereby to define a cell first end wall.

mounting a second thin electrode layer to the other side of the sheet and extending over the aperture whereby to define a second cell end wall in substantial

10 overlying registration with the first electrode, and

providing means for admission of a liquid into the cell defined between the side wall and said end walls.

2. A method according to claim 1 wherein at least one of the electrode layers is a noble metal.

15 3. A method according to claim 2 wherein at least one of the electrode layers is palladium.

4. A method according to any one of claims 1 to 3 wherein each of the metal electrode layers substantially covers the aperture.

5. A method according to any one of the preceding claims wherein the aperture is 20 of circular cross-section, whereby the side-wall is cylindrical.

6. A method according to any one of the preceding claims wherein the means for admission of a liquid comprise a slot or passage communicating between the cell interior and an edge of the sheet.

7. A method according to any one the preceding claims wherein the thin electrode is a sputter coated metal deposit.
8. A method according to any one of the preceding claims wherein the electrode layers are adhered to the sheet.
- 5 9. A method of manufacture of an electrochemical cell comprising the steps of:
 - depositing a first sputter coating of a metal on a first suitable substrate,
 - depositing a second sputter coating of the same or a different metal on a second suitable substrate,
 - forming an aperture in a sheet of electrically resistive material, said aperture
- 10 defining a side wall of the cell and having openings on each face of the sheet
 - adhering the first substrate to the sheet with the metal coating thereof adjacent one side of the sheet whereby to expose a portion of the first coating to one aperture opening and to provide a first electrode
 - adhering the second substrate to the sheet with the metal coating thereof
- 15 adjacent the other side of the sheet whereby to expose a portion of the second coating to the other aperture opening facing the first and to provide a second electrode, and
 - providing means for admission of a liquid into the cell defined between the side wall and said exposed portions of metal coating.
10. A method according to claim 9 wherein the means for admission of a liquid
- 20 comprise a passage communicating between the cell interior and an edge of the sheet.
11. A method according to any one of claims 9 or 10 wherein the first sputter coating and the second sputter coating are adhered to the sheet by means of an adhesive.
12. A method according to claim 11 wherein the adhesive is a heat activated adhesive.

13. A method according to any one of the preceding claims wherein one or more chemicals for use in the cell are printed onto one or both electrodes.
14. A method according to any one of the preceding claims wherein one or more chemicals for use in the cell are added in a solvent to the cell after one electrode has been
5 adhered to the sheet and before the other electrode has been adhered to the sheet and wherein the solvent is subsequently evaporated to leave the chemicals in dry form in the cell.
15. A method according to any one of the preceding claims wherein one or more chemicals for use in the cell are contained in or on a support which is included within the
10 cell.
16. A method according to any one of the preceding claims wherein one or more chemicals for use in the cell are contained within the cell in self-supporting form.
17. A method according to any one of the preceding claims wherein one or both electrodes comprise a sputter coated metal and one or more chemicals for use in the cell
15 are printed onto the metal surface immediately after the metal is sputter coated.
18. A method according to claim 17 wherein one or more chemicals for use in the cell are printed onto the metal surface by means of an ink jet printer.
19. A method according to any one of the preceding claims wherein a plurality of first sputter coatings of metal are formed on a first substrate, a plurality of second sputter
20 coatings of metal are deposited on a second substrate, a plurality of apertures are defined in a sheet of electrically resistant material, each aperture defining a side wall of a cell, laminating the first substrate to the sheet with the metal coatings thereof adjacent one side of the sheet and defining first end walls of respective cells defined by the sheet, and adhering the second substrate to the sheet with the metal coatings thereof adjacent the

other side of the sheet and defining second cell end walls of respective cells defined in the sheet, and subsequently dividing the assembly into a plurality of cells.

20. A method according to any one of the preceding claims, wherein one or more of the parts is provided with a formation adapted to interlock with a formation of another 5 one of the parts or of an assembly jig.

21. A method according to any one of the preceding claims, wherein the sheet of electrically resistant material comprises a laminate having two layers, each of electrically and chemically resistant material, having a metal layer therebetween which in the final product is useful as a reference electrode disposed intermediate the first metal and the 10 second metal.

22. An electrochemical cell comprising an electrically resistive substrate having a first thin layer of first metal on one face, a second electrically resistive substrate having a second thin layer of metal sputter coated on the other face. said substrates being disposed with the metal coating of one facing the metal coating of the other and being separated 15 by a sheet pierced by an aperture, the wall of which aperture cooperates with said metal coatings to define a cell wall, and an aperture whereby a solution may be introduced into the cell.

23. A method according to any one of the preceding claims, wherein the substrate is a Polyethylene Terephthalate.

20 24. A method according to any one of the preceding claims, wherein the sheet is of Polyethylene Terephthalate.

25. A method according to any one of the preceding claims, wherein at least one of the electrode layers comprises palladium.

- 16 -

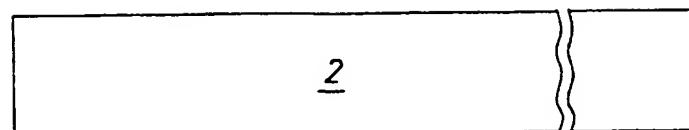
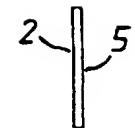
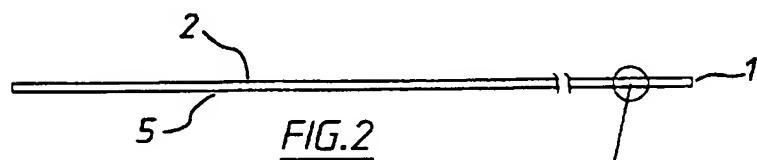
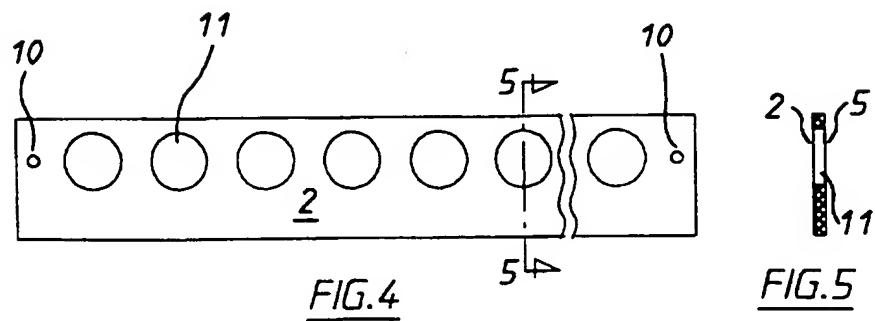
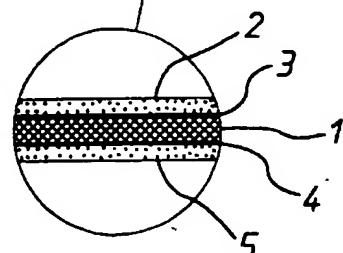
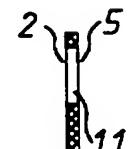
26. A method according to any one of the preceding claims, wherein one of the sputtered metal layers is of silver.
27. A method according to any one of the preceding claims when one of the layers is a metal coating of silver and is halogenised to produce a silver halide surface.
- 5 28. A method according to any one of the preceding claims further comprising a second aperture piercing the sheet, and wherein the second aperture is covered on at least one side of the sheet by the same metal layer as covers the first aperture on that side.
29. A method according to anyone of claims 1 to 27 further comprising a second aperture wherein the second aperture is covered on at least one side by a metal layer
- 10 which is electrically insulated from the metal layer covering the first aperture on that side.
30. A method according to claim 1, further comprising the step of providing means for venting of gas during the admission of liquid into the cell.
31. A method according to claim 6, wherein the slot or passage is formed through
- 15 the side wall and through the sheet of electrically resistive material.
32. A method according to any one of the preceding claims wherein the second thin electrode layer is mounted in opposing relationship a distance of less than 200 microns from the first thin electrode layer.
33. A thin layer electrochemical cell comprising
- 20 an electrically resistive sheet pierced by an aperture,
- a first electrode layer covering the aperture on one side of the sheet,
- a second electrode layer covering the aperture on the other side of the sheet, and
- a passage for admission into the aperture of a liquid.

34. A cell according to claim 33 wherein at least one electrode layer comprises a conductor.
35. A cell according to claims 33 or 34 wherein at least one electrode layer comprises a semi-conductor.
- 5 36. A cell according to any one of claims 33 to 35 wherein at least one electrode layer comprises a noble metal.
37. A cell according to any one of claims 33 to 36 wherein at least one electrode coating is palladium.
- 10 38. A cell according to any one of claims 33 or 37 wherein an electrode comprises a sputter coated metal.
39. A cell according to any one of claims 33 to 38 wherein the cell contains one or more reagents.
40. A cell according to claim 33 wherein at least one reagent is provided on a sputter coated metal electrode.
- 15 41. A cell according to claims 33 or 40 wherein one or more reagents are supported or impregnated in a carrier within the cell.
42. A cell according to any one of claims 33 to 41 wherein one or more chemicals for use in the cell are contained within the cell in self-supporting form.
43. A cell according to any one of claims 33 to 42 wherein at least one of the electrodes is a sputter coated palladium.
- 20 44. A cell according to any one of claims 33 to 43 wherein the electrodes are of substantially the same area.
45. A cell according to any one of claims 33 to 44 wherein the electrodes are in substantial registration one with the other.

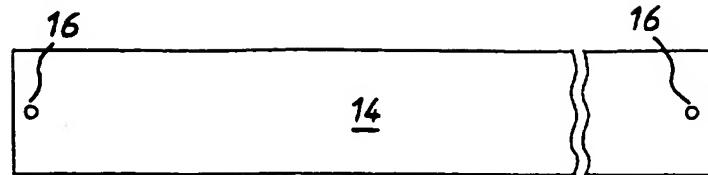
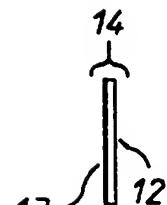
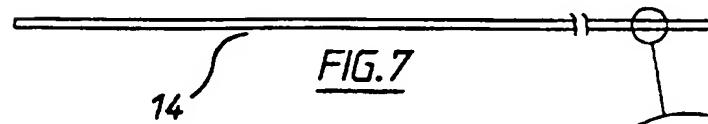
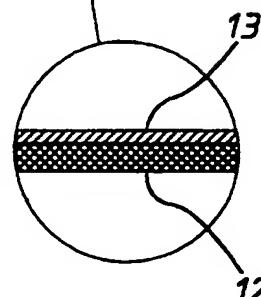
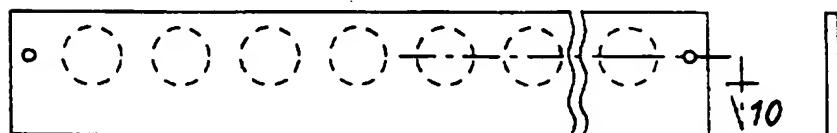
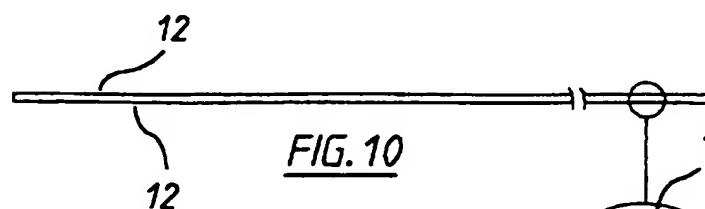
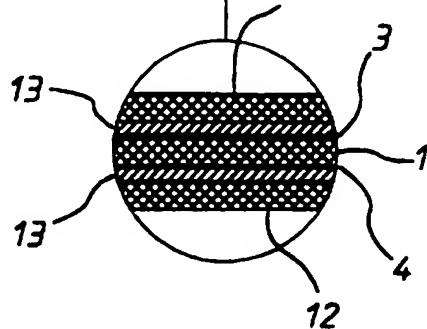
- 18 -

46. A cell according to claim 33 further comprising means for venting of gas during admission of liquid into the cell.
47. A cell according to any one of claims 33 to 46 wherein the second electrode layer is mounted in opposing relationship a distance of less than 200 microns from the first electrode layer.
5

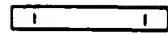
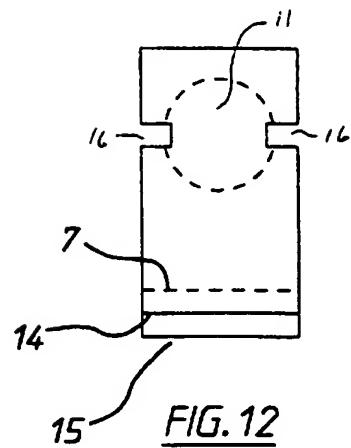
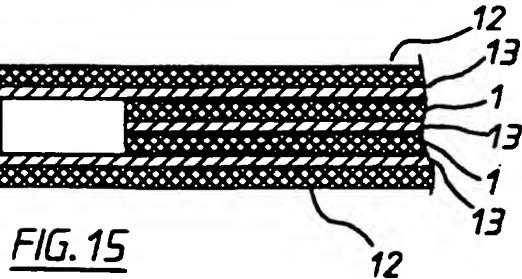
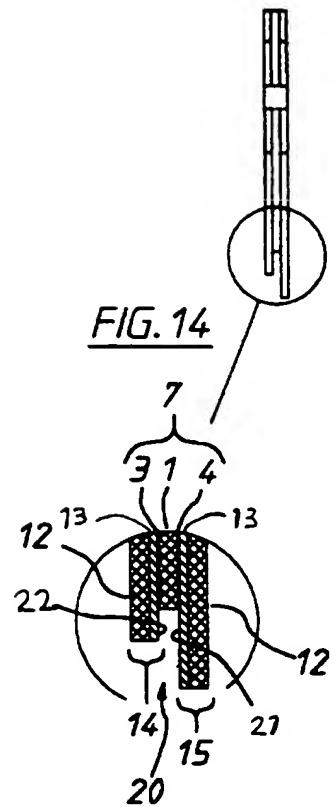
1 / 3

FIG. 1FIG. 3FIG. 2FIG. 4FIG. 5

2/3

FIG. 6FIG. 8FIG. 7FIG. 11FIG. 9FIG. 10

3 / 3

FIG. 13FIG. 14

INTERNATIONAL SEARCH REPORT

International Application No.
PCT/AU 96/00724

A. CLASSIFICATION OF SUBJECT MATTER

Int Cl⁶: G01N 27/403

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC : G01N 27/40, 27/403, 27/42, 27/49, 27/52

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
AU : IPC as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
WPAT, JAPIO

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GB 2 235 050 A (Sieger Limited) 20 February 1991 entire document, Figure 1	1-6, 25, 33, 34, 37, 39, 44, 45
Y	AU 54873/94 A (ASULAB S.A.) 11 August 1994 entire document	2, 7, 25-27, 36-38, 43
Y	US 5 393 399 (Van den BERG et al.) 28 February 1995 entire document	2, 27, 36



Further documents are listed in the continuation of Box C



See patent family annex

* Special categories of cited documents:	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E" earlier document but published on or after the international filing date	"Y"	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&"	document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means		
"P" document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search 20 December 1996	Date of mailing of the international search report 09.01.97
Name and mailing address of the ISA/AU AUSTRALIAN INDUSTRIAL PROPERTY ORGANISATION PO BOX 200 WODEN ACT 2606 AUSTRALIA Facsimile No.: (06) 285 3929	Authorized officer A. ALI Telephone No.: (06) 283 2607

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/AU 96/00724

C (Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	Patent Abstracts of Japan, C-484, page 110, JP 62-228 274 A (MITSUBISHI CHEM IND LTD) 7 October 1987	2, 27, 36
A	EP 299 779 A (SRI INTERNATIONAL) 18 January 1989 entire document	

INTERNATIONAL SEARCH REPORT
Information on patent family members

International Application No.
PCT/AU 96/00724

This Annex lists the known "A" publication level patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent Document Cited in Search Report				Patent Family Member		
US	5393399	CA	2105510	EP	586982	FR
		JP	6213863			2695481
AU	94/54873	CA	2114685	EP	609760	FR
		JP	6294769	US	5395504	2701117
GB	2235050	DD	297251	DE	4025635	
EP	299779	CA	1291208	JP	1088245	US
JP	62228274					4812221
END OF ANNEX						